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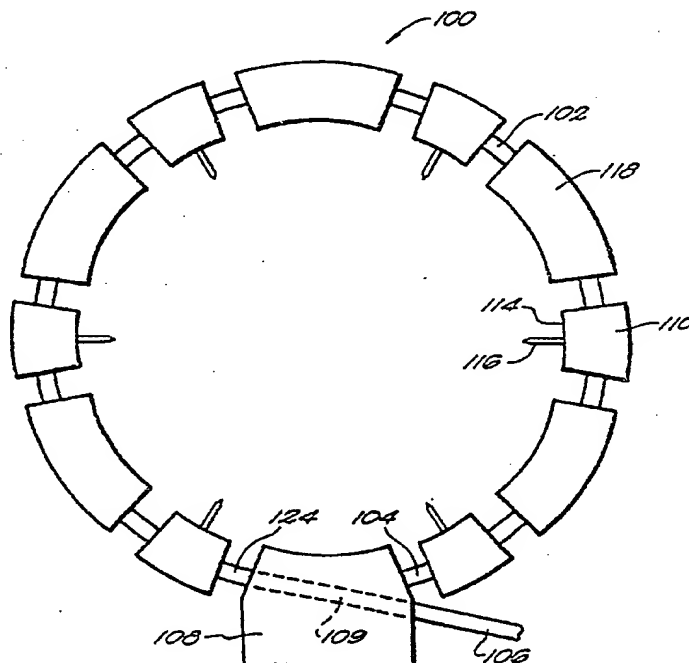
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(54) Title: EXOVASCULAR ANASTOMOTIC DEVICE

(57) Abstract

An anastomotic device (100) includes a flexible member (102), formable into a loop and a plurality of tissue penetrating anchoring elements (110) slide mounted on the flexible member. The tissue penetrating anchoring elements (116) extend generally towards a center of the loop formed by the flexible member. The device is adapted for anastomosing vessel segments or a vessel and a graft. In a further embodiment, an exo-vascular anastomotic device is suitable for end to side anastomoses, and it includes an anastomotic ring and a harness assembly.



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5 EXOVASCULAR ANASTOMOTIC DEVICE

FIELD OF THE INVENTION

The present invention relates to devices useful with surgical anastomoses and more particularly to a sutureless anastomotic device.

10 BACKGROUND OF THE INVENTION

The joining of one hollow or tubular organ to another in a patient, known as anastomosis, is a common surgical procedure. Blood vessels serve as an example of such organs for which anastomoses may be required. For example, a sharp foreign object or severe trauma can sever a blood vessel, requiring an anastomotic
15 procedure to restore patency of the vessel. If a vessel is occluded or otherwise diseased, a bypass procedure may include replacing the occluded section with a portion of a vessel from another part of the body. Also, if a portion of a vessel is diseased or damaged, a synthetic graft, or prosthesis, may be introduced to replace that portion of the vessel.

Anastomoses procedures have been traditionally effected by using sutures to
20 join the vessels and/or grafts. It will be appreciated by one of ordinary skill in the art that there are significant disadvantages associated with suturing vessels. For example, a high level of skill is required to properly join the vessels and/or grafts. The suturing of vessels and/or grafts is, moreover, a procedure that is quite time
25 consuming. To anastomose a blood vessel having a diameter in the range of about five to twenty-five millimeters, approximately twenty to thirty well placed stitches must be placed about the circumference of the vessel, requiring at least fifteen minutes under optimal conditions.

The suturing process is further complicated by friable and rigid, calcified
30 vessels increasing the time required for the procedure. As is known to one of ordinary skill in the art, longer operative times are associated with a greater likelihood of heart attack and infection. Moreover, any improper suturing can result in serious medical consequences, including hemorrhaging and infection.

Anastomosis of living vessels requires the vessel layers to be positioned
35 properly with respect to each other. As shown in FIG. 1, a vessel, such as a human

5 artery 10 has an inner layer (intima) 12, a middle layer (media) 14, and an outer
layer (adventitia) 16. To anastomose a vessel or graft, the sutures must be precisely
placed using a curved needle to pierce the vessel first from the outside, and then
from the inside back to the outside of the vessel. As noted above, such a suturing
10 process is time consuming and demands a high level of precision. Time is
especially critical in procedures that require repair of a major artery.

Suture-based anastomoses have inherent limitations because the mechanical
strength of the sutures join the graft and the vessel. Suturing can be especially
difficult in cases where a vessel is so diseased or so severely damaged that there is
only a limited amount of healthy tissue to which the sutures can be anchored. The
15 brittleness of some diseased vessels can render them less amenable to suture-based
anastomotic procedures.

There are inherent advantages to minimally invasive laparoscopic procedures.
However, the limited degree of freedom afforded by the closed surgical environment
compounds the difficulties associated with suture-based anastomotic procedures.

20 Because suture-based anastomotic procedures suffer from the drawbacks
noted above, attempts have been made to develop sutureless anastomotic devices.
See, for example, U.S. Patents Nos. 4,470,415, 4,917,087, 4,930,674, and
5,486,187. However, the complexity of such devices and/or their lack of ease of
use have limited their clinical acceptance. There is thus a need for a sutureless
25 anastomotic device that can be easily and effectively used in a variety of anastomotic
procedures.

SUMMARY OF THE INVENTION

30 The present invention provides a sutureless anastomotic device useful in
sutureless anastomotic procedures. In one embodiment the anastomotic device is an
exovascular anastomotic device. Although the invention is primarily described and
illustrated as an exovascular anastomotic device, it is understood that it can be used
for other anastomoses procedures such as gastro-intestinal anastomoses.

35 The device, in its initial configuration, is an elongate flexible member having
first and second ends. One or more anchoring elements are disposed on the flexible

5 member. The anchoring elements preferably are each in the form of a tubular body having an axial passageway formed therein. One surface of the anchoring element has formed thereon at least one tissue-penetrating projection. The anchoring elements are preferably slidably mounted, through the axial passageway, upon the flexible member. Optimally, one or more spacer elements may be disposed on the flexible member between each, or selected, adjacent anchoring members to control the positioning of the anchoring members.

10 The flexible member is formable into a loop of an adjustable diameter. In one embodiment, one of the first or second ends of the flexible member includes a locking mechanism having an opening formed therein. In operation, the other of the first or second ends of the flexible member is threaded through the opening of the locking mechanism so that the flexible member assumes the shape of a loop. In this loop configuration, the anchoring elements are circumferentially oriented with the tissue-penetrating projections being inwardly directed, either radially inward or at a selected angle to the radius. Preferably, the anchoring elements are slidably mounted on the loop so that before installation of the device in an anastomotic procedure they can be positioned as needed; once the device is initially placed on a vessel the anchoring elements are stationary.

15 Surface features are preferably formed on at least one surface of the flexible member. These surface features interact with a complementary surface feature or other structure formed in the opening of the locking mechanism so that the end inserted into the locking mechanism can only be moved in one direction with respect to the locking mechanism. Preferably, this unidirectional movement enables the loop to be adjusted only in a manner to decrease its diameter.

20 In one embodiment, the anastomotic device is used by forming the flexible member into a loop that is circumscribed about an end of a first vessel segment. The end of the first vessel segment is then overlapped with an end of a second vessel segment. An initial adjustment to the diameter of the loop is made to cause the tissue-penetrating projections to penetrate the vessels. Thereafter, the flexible member is moved with respect to the anchoring elements (which remain stationary) to reduce the size of the loop until it is properly dimensioned about the vessels to

5 anastomose the first and second vessel segments. An intravascular obturator (such as an obturator balloon) may be positioned within the vessel to assist in the anastomotic procedure.

In another embodiment, the anastomotic device of the invention can be used for an end-to-side anastomosis. In this embodiment, the device includes an
10 anastomotic ring and a harness assembly having a harness ring coupled to at least one locking band. The anastomotic ring optionally includes at least one filamentary member fixedly secured to the anastomotic ring.

An end-to-side anastomosis can be conducted by placing a vessel or graft within the harness ring and then the anastomotic ring, and everting an end of the
15 vessel or graft segment over the anastomotic ring. A receiving vessel is cut to form a substantially round hole, the opening of which tapers inwardly from the adventitia to the intima. The everted end of the vessel or graft is placed into the complementary hole formed in the receiving vessel and the harness ring is urged toward the anastomotic ring by cinching the locking band to the receiving vessel.
20 The harness ring retains the everted end of the graft and anastomotic ring in the hole in the receiving vessel. The filamentary member(s) can secure the anastomotic ring to the harness assembly.

BRIEF DESCRIPTION OF THE DRAWINGS

25 The invention will be more fully understood from the following detailed description taken in conjunction with the following drawings, in which:

FIG. 1 is a prior art perspective view of an artery;

FIG. 2 is a side view of an exovascular anastomotic device in accordance with the present invention shown in an unengaged position;

30 FIG. 3 is a side view of the anastomotic device of FIG. 2 shown in an engaged position;

FIG. 4 is a side view of an anchoring element forming a portion of the anastomotic device of FIG. 2;

35 FIG. 5 is a enlarged sectional view of a portion of the locking mechanism and the flexible member, forming a portion of the anastomotic device of FIG. 2;

5 FIG. 6 is a sectional view of a spacer element forming a portion of the anastomotic device of FIG. 2;

 FIG. 7 is a cross sectional view of the anastomotic device of FIG. 2 disposed on a vessel;

 FIG. 7A is a perspective view of the anastomotic device of FIG. 7;

10 FIG. 7B is an enlarged cross sectional view of an alternative embodiment of the anastomotic device of FIG. 7;

 FIG. 7C is a side view of a further embodiment of the anastomotic device of FIG. 7;

15 FIG. 7D is a perspective sectional view of an obturator adapted for use in conjunction with the anastomotic device of FIG. 7;

 FIG. 8 is a cross sectional view of an alternative embodiment of an anastomotic device in accordance with the present invention;

 FIG. 9 is a cross sectional view of a further embodiment of an anastomotic device in accordance with the present invention;

20 FIG. 9A is a perspective view of a prior art vessel having a hole punched therein;

 FIG. 9B is a cross sectional view of the vessel of FIG 9A;

 FIG. 10 is a perspective view of an anastomotic ring forming a portion of the anastomotic device of FIG. 9; and

25 FIG. 11 is a top view of a harness assembly forming a portion of the anastomotic device of FIG. 9.

DETAILED DESCRIPTION OF THE INVENTION

30 The drawings are understood to be illustrative of the concepts disclosed herein to facilitate comprehension of the invention. Further, the drawings are not to scale, and the scope of the invention is not to be limited to the particular embodiments shown and described herein.

35 Referring to FIGS. 2-8, a sutureless anastomotic device 100 includes a flexible member 102 having a first end 104 and a second end 106. A locking mechanism 108, or similar device, enables the formation of a closable loop and is

5 secured to the first end 104 of the flexible member 102. The second end 106 is threaded through an opening 109 in the locking mechanism to form a loop. Anchoring elements 110 are slidably mounted on the flexible member 102. The anchoring elements 110 are preferably in the form of a tubular body 111 defining an axial passageway 112 therein. The anchoring elements 110 each include a first
10 surface 114 having formed thereon a tissue-penetrating projection 116. In an exemplary embodiment, one or more spacer elements 118 can be disposed on the flexible member 102 between adjacent anchoring elements 110 to control the positioning and/or spacing of the anchoring elements.

15 It is understood that various alternative structures may be used as an alternative to locking mechanism 108. Suitable structures must allow the formation of a loop, having an adjustable diameter, from an initially elongate member. For example, the locking mechanism can include a screw element, advancement of which moves one end of the flexible member relative to the other end of the flexible member to reduce the diameter of the loop.

20 Each of the tissue-penetrating protrusions 116 include a first end 119 coupled to the tubular body, a generally pointed second end 120, and an intermediate portion 122. In an exemplary embodiment, the intermediate portion 122 has an annular cross section defining a substantially constant diameter along the length thereof. The tissue-penetrating protrusion 116 has a length in the range of about 0.1 to 5
25 millimeters, and preferably is about one millimeter. The diameter of protrusion 116 at the intermediate portion 122 is in the range of about 0.01 to 0.3 millimeters, and preferably about 0.1 millimeters. Other geometries for the tissue-penetrating protrusions 116 are also possible, such as conical and flattened, for example.

30 The anchoring elements 110 are formed from biocompatible materials having good mechanical properties. Exemplary materials include stainless steel, titanium, titanium alloys, metals, ceramics, and suitably rigid, biocompatible polymers. The device 100 is shown having six anchoring members 110, each having one tissue-penetrating protrusion 116. However, it is understood that a greater or lesser number of anchoring members 110 may be used, and that a
35 member 110 may have more than one protrusion 116 (FIG. 7C). In a further

5 embodiment, the second end 120 of the protrusions includes one or more barbs 123 (FIG. 7B) for securing the protrusions in tissue. Further, the protrusions 116 can extend from the tubular body 111 at any point on the first surface 114.

10 In an exemplary embodiment, the anchoring elements 110 are freely movable with respect to the flexible member 102 before protrusions 116 are embedded in tissue. The flexible member 102 is of sufficient width so as to provide lateral stability for the tissue-penetrating protrusions 116. Generally, the width of flexible member 102 ranges from about 1.5 millimeters to 2 centimeters, and more preferably from about 3 to 4 millimeters. The tissue-penetrating protrusions 116 should generally be oriented to form an acute angle with respect to a radial center point of the loop (FIG. 7B). This angle is generally in the range of about five to
15 eighty degrees, and more preferably is about thirty degrees. The optimal angle for the protrusions 116 is determined by the amount of deformation that occurs as the protrusions penetrate tissue. More specifically, the more rigid the protrusions 116, the more the protrusions can be angled with respect to the radial centerpoint of the loop. In another embodiment, the protrusions 116 are directed to the radial
20 centerpoint of the loop formed by the flexible member.

As illustrated in Figures 2 and 5, the flexible member 102 defines a first surface 124 having a plurality of surface features 126, which in an exemplary embodiment are spaced teeth. The surface features 126 can be disposed on the first
25 surface 124 or an opposing surface of the flexible member 102, i.e., inward or outward with respect to the loop. The opening 109 in the locking mechanism 108 may include a structure such as a surface feature in the form of an arm 127, or a similar member, that is engageable with a respective one or more of the teeth 126 associated with the flexible member 102. In a preferred embodiment, each of the
30 teeth 126 includes a sloped surface 128 defining an acute angle with respect to the longitudinal axis X of the flexible member and an engagement surface 130 extending generally perpendicular to the longitudinal axis X. It will be appreciated by one of ordinary skill in the art that the size of the teeth 126 may vary substantially. In an exemplary embodiment, the sloped surface 128 is about 0.3 millimeters in length
35 and the engagement surface 130 is about 0.1 millimeters in height. The arm 127 is

5 biased towards the flexible member 102 and contacts the engagement surface 130 of
the respective tooth. As the flexible member 102 is moved in a first direction,
indicated by arrow 132, the arm 127 elastically deforms as it slides up the sloped
surface 128 and then returns to its original position as it reaches an adjacent
10 engagement surface. The arm 127 operates in conjunction with the engagement
surface 130 to prevent movement of the flexible member 102 in a direction opposite
of the first direction 132. Thus, the surface features of the locking mechanism and
the flexible member permit unidirectional movement of the flexible member with
respect to the locking mechanism 108 to allow only a reduction in the size of the
loop.

15 The flexible member 102 can be made from a variety of suitably flexible
materials including biocompatible polymers. Exemplary materials include nylon
polymers. The flexible member 102 can be formed from inelastic or moderately
elastic materials known to one of ordinary skill in the art. Moderately elastic
flexible members are useful as they expand in response to a pulse of a fluid through
20 a vessel. In a further embodiment useful in gastro-intestinal applications, the device
is formed from biodegradable polymers, such as polyglyconate, so that the device
need not be surgically removed after healing has taken place. Commercially
available examples of such materials include POLYSORB and MAXON.

25 As shown in Figure 6, the optional spacer elements 118 preferably are
tubular in shape, defining an aperture 134 through which the flexible member 102
may pass. Each of the spacer elements 118 includes an outer surface 136 and an
inner surface 138 (as configured on the loop), wherein the inner surface is about the
same distance from the flexible member 102 as the surface 114 of the anchoring
elements 114. In one embodiment, one or more spacer elements 118 are located
30 between adjacent tissue-penetrating protrusions 116 so that the anchoring elements
are generally equally spaced about the loop as the size of the loop is reduced.
However, it is understood that the anchoring elements need not necessarily be
equally spaced. The spacer elements 118 preferably have sufficient flexibility or
other physical characteristics (e.g., geometry) to enable them to conform to the
35 curvature of the loop.

5 The dimensions of the spacer element can vary depending upon the requirements of a given application. The thickness (or height) of the spacer elements should be consistent with the thickness of the tubular body portion of anchoring elements 114 to form a substantially continuous contact surface with a vessel or a graft. Typically, the thickness is in the range of about 0.02 to 0.2
10 millimeters. The width of the spacer elements 118 is in the range of about one to twenty millimeters, and preferably, is about the same width as the anchoring elements. The length of spacer elements 118 is in the range of about one half to about twice the length of the anchor elements, and it is understood that the spacer elements can be longitudinally compressible.

15 In one embodiment, the spacer elements can have a longitudinal slit along the length thereof so that one or more selected spacer elements can be removed or added to the flexible member 102. In addition, a series of spacer elements 118 of the same or differing lengths can be disposed between adjacent anchor elements 110 to provide a desired spacing of the anchoring elements (FIG. 7C). One or more
20 selected spacer elements can be removed or added to adjust the minimum diameter of the loop. In another embodiment, the spacer elements are formed from a material that can be cut with relative ease so that one or more selected spacer elements can removed from the loop.

25 One of ordinary skill in the art will appreciate that the spacer elements can be made from a variety of biocompatible materials, including metals, polymers and ceramics. Polymeric materials are preferred for most applications, and suitable exemplary polymers include polytetrafluoroethylene (PTFE) and other polymers known to those of ordinary skill in the art.

30 Figures 7 and 7A illustrate the use of a sutureless anastomotic device 100 according to the present invention in a distal anastomosis of an end 150 of a native blood vessel 152 and an end 154 of a graft 156. As indicated by arrow 159, the blood flow is in the direction from the graft 156 to the vessel 152. To effect this procedure, the end 154 of the graft is moved toward the device 100 so that the device, in the form of a loop, surrounds the graft 156. The end 150 of the vessel
35 152 is inserted within the end 154 of the graft so that portions of the vessel and the

5 graft overlap at an overlap area 158. The flexible member 102 of the device is moved with respect to the locking mechanism (and the anchoring members) to decrease the size of the loop. As the loop contracts, the tissue-penetrating protrusions 116 penetrate the graft 156 and then the vessel 152.

10 Contemporaneously, the spacer elements 118 slide with respect to the flexible member 102 so that surfaces 114,138 of the anchoring elements and the spacer elements attain circumscribed engagement with an exterior surface 160 of the graft. As noted above, the anchoring elements 110 are freely slidable upon the flexible member 102. Once protrusions 116 penetrate tissue, however, anchoring elements 110 are stationary with respect to flexible member 102.

15 Upon initial penetration of the surface of the graft 156, the tissue-penetrating protrusions 116 are urged radially inward by the flexible member 102, without circumferential movement, as the loop diameter is further decreased. If the protrusions 116 are angled with respect to a radial centerpoint of the loop, tangential and radial force vectors are applied to the protrusions as the loop size decreases.

20 The spacer elements 118 and the anchor elements provide substantially continuous contact with and substantially constant pressure on the graft outer surface 160 to prevent gaps that could result in leakage of blood between the artery 152 and the graft 152. The device is of sufficient flexibility so as to conform to any irregularities in the outer surface of the vessel.

25 The diameter of the loop is decreased by moving the flexible member 102 with respect to the locking mechanism 108 and the anchoring elements. One of ordinary skill in the art will appreciate that a variety of suitable applicator devices can be used. In one example, a suitable applicator gun (not shown) includes a head portion, a manual actuator mechanism, and a handle. In operation, the head portion abuts the locking mechanism 108 and the actuator mechanism engages the surface features 124 of the flexible member 102 for longitudinally displacing the flexible member with respect to the locking mechanism. A user, such as a surgeon, activates the actuator mechanism to manipulate the loop to a desired size.

30

35 In another embodiment, shown in Figure 7C, the anastomotic device 100 can include one or more crimping elements 155 movably disposed on the flexible

5 member 102. Such crimping members can be useful to close any gap 161 formed
between the device 100 and an external surface of a graft. The crimping element
155 includes a tubular body 157 and an arcuate projecting member 159 extending
from the tubular body. The projecting member 159 substantially conforms to the
10 arcuate shape of the graft, thereby eliminating any gap. In addition, where a graft
is to be anastomosed to a vessel having a smaller diameter, any excess graft material
can be gathered at an end 163 of the projecting member 159 and pressed against an
adjacent spacer element 118 or anchor element 110 to eliminate spaces or wrinkles
in the graft.

15 In one embodiment, illustrated in Figure 7, the tissue-penetrating protrusions
116 are of a length such that they do not extend into the lumen of the affected
vessel. Instead, the tissue-penetrating protrusions 116 lodge in the healthy adventitia
of the vessel, and do not affect any diseased intima having a layer of calcified
plaque 162. A balloon or rigid obturator (e.g., 181 in FIG. 7D) can be inserted into
20 the lumen of the vessel in a manner known to those of ordinary skill in the art to
prevent collapse of the vessel as the device 100 is actuated. In one example, the
obturator can be inserted by way of a catheter through the groin area.

As shown in FIG. 7D, a balloon obturator 181 can include an inner layer 183
and a sheath 185 surrounding the inner layer. The sheath 185 resists penetration by
the tissue-penetrating protrusions 116 and protects the inner layer 183 from puncture
25 as the anastomotic device anastomoses one vessel to another. The sheath 185 is
formed from suitable puncture resistant materials, such as latex.

30 In another embodiment, shown in FIG. 8, a device 100' includes tissue-
penetrating protrusions 116' having a length sufficient to penetrate the intima and
extend into the lumen of a vessel and into contact with a rigid obturator 164. Such
contact with obturator 164 causes the tissue-penetrating protrusions 116' to deform
as the device fully engages the vessel. The resulting deformed tissue-penetrating
protrusions 116' can help to maintain a seal between the graft 156 and vessel 152 to
prevent or minimize any leakage of blood between the vessel and the graft. This
embodiment is well suited for a distal anastomosis in which the graft overlaps the
35 end of the vessel and blood flows from the graft to the vessel.

5 Figures 9-11 illustrate an anastomotic device 200 that is useful for an end-to-
side anastomosis. Device 200 includes a semi-rigid anastomotic ring 202 and a
harness assembly 204. The harness assembly 204 includes a semi-rigid harness ring
206 and a pair of flexible members 208 secured thereto. The harness assembly 204
10 can include connecting portions 209 extending laterally from the flexible members
208 to the harness ring 206. The anastomotic ring 202 includes a tapered outer
surface 210 and, optionally at least one filamentary member 212 extending from the
anastomotic ring 202 that can be used to secure the anastomotic ring to the harness
assembly 204. The harness ring 206 includes a radially overlapping portion with
15 respect to the anastomotic ring 202 as the inner diameter of the harness ring is less
than the outer diameter of the anastomotic ring.

 In operation, device 200 is used by first placing a first vessel segment 214
through the harness ring 206 and the anastomotic ring 202. A portion 216 of the
first vessel segment 214 is everted over the anastomotic ring 202. A hole 217 is
20 then cut in a second vessel 218 by means of a puncher using techniques well known
to those of ordinary skill in the art (FIGS. 9A, 9B). The hole 217 has a surface 219
that is tapered inwardly from the adventitia to the intima. The hole 217 has a
diameter slightly smaller than the diameter of the anastomotic ring 202.

 The everted portion 216 is then placed into the hole formed in the second
vessel 218. The harness assembly 204 is cinched down to the second vessel 218 by
25 the flexible members 208 in conjunction with locking mechanisms 220. As the
harness assembly 204 is secured to the second vessel, the overlapping portion of the
harness ring 206 and anastomotic ring 202 retains the everted portion 216 of the
first vessel in sealed engagement in the hole in the second vessel. The optional
filamentary members 212 can then be used to secure the anastomotic ring to the
30 harness assembly 204 and vessel and the connecting portions 209 cause the harness
and anastomotic rings 206,202 to conform to the annular outer surface of the second
vessel 218.

 For a substantially perpendicular end-to-side anastomosis of the first and
second vessels 214,218, the hole 217 and corresponding harness and anastomotic
35 rings 206,202 will be annular. If the first vessel 214 is to be anastomosed at an

5 angle with respect to the second vessel 218, the formed hole 217 and corresponding rings 206,202 will be generally oval in shape.

10 The anastomotic junction of the first and second vessels 214,218 must not leak in order to achieve a successful anastomosis. The anastomotic device 200 can include additional features to assist in sealing the vessels. For example, a type of
15 bio-sealant 211, of the type known to one of ordinary skill in the art, can be placed under and in the area of the harness ring 206 to seal the harness ring and the anastomotic ring 202 and/or first vessel. Also, the harness assembly 204 can include tissue-penetrating teeth 225, which can be barbed, to secure and seal the harness assembly and the second vessel 218. In addition, the harness ring 206 can
20 include a surface feature, such as a ridge, to enhance sealing engagement with a surface feature in the opposing surface of the anastomotic ring 202. Various complementary surface features on the harness assembly 204 and the anastomotic ring 202 can be also be used. A further sealing enhancement utilizes one or more surgical clips for grasping a portion of the first vessel 214, the harness ring 206, and/or the second vessel 218. In a still further embodiment, the portion of the vessel everted over the anastomotic ring 202 can extend further so as to be engaged by the harness ring. The harness and anastomotic rings 206,202 can include one or more complementary surface features 221 that can penetrate through the everted portion of the first vessel and secure the harness and anastomotic rings together.

25 The sutureless anastomotic device of the present invention can be used both in open surgery and in closed surgery, such as laparoscopic surgery. Such closed surgical techniques provide significant advantages over more invasive, open surgical techniques where much larger incisions are required. The benefits of closed surgery include reduced recovery time, decreased pain or discomfort and a considerable
30 cost savings. Another important advantage of the sutureless anastomotic device of the present invention is the reduced time required to anastomose vessels as compared with known suturing techniques. This reduction in time significantly reduces risk to the patient and increases the success rate of surgical procedures.

35 The exovascular device provides further advantages over sutures. The device can be anchored to healthy adventitia whereas sutures must penetrate plaque layered

on the intima of the vessel, risking dislodgement of chunks of plaque.

One of ordinary skill in the art will realize further features and advantages of the invention from the above-described embodiments. Accordingly, the invention is not to be limited by what has been particularly shown and described, except as indicated by the appended claims. All references cited herein are expressly incorporated by reference in their entirety.

What is claimed is:

- 5 1. An anastomotic device, comprising:
 a flexible member having a first end and a second end and first and second
surfaces, the flexible member being formable into a loop of an adjustable size; and
 one or more anchoring elements mounted upon the flexible member, each
0 anchoring element having at least one tissue penetrating protrusion formed on one
surface thereof.
2. The anastomotic device according to claim 1, wherein at least one of the
anchoring elements is slidably mountable upon the flexible member.
- 5 3. The anastomotic device according to claim 1, further including a locking
mechanism disposed on one of the first or second ends of the flexible member, the
locking mechanism including an opening formed therein for receiving the other of
the first or second ends of the flexible member.
- 0 4. The anastomotic device according to claim 3, wherein the locking mechanism
is immovably affixed to the first end of flexible member.
5. The anastomotic device according to claim 3, wherein the opening of the
locking mechanism includes at least one surface feature and one of the first or
15 second surfaces of the flexible member includes a plurality of surface features, the
at least one surface feature of the locking mechanism being engageable with a
respective one or more of the surface features on the flexible member.
6. The anastomotic device according to claim 5, wherein the at least one surface
30 feature of the locking mechanism and the plurality of surface features of the flexible
member allow only unidirectional movement of the flexible member with respect to
the locking mechanism.

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5 7. The anastomotic device according to claim 6, wherein the surface features of the flexible member are mounted on the first surface of the flexible member, and comprise teeth-like members having a first surface ramped with respect to a longitudinal axis of the flexible member and a second surface orthogonal with respect to the longitudinal axis of the flexible member.

10 8. The anastomotic device according the claim 7, wherein the teeth-like surface features of the flexible member permit only unidirectional movement of the flexible member within the locking mechanism, the unidirectional movement taking place in direction to enable only a decrease in the diameter of the loop.

15 9. The anastomotic device according to claim 1, wherein the at least one tissue-penetrating protrusion of the anchoring member is oriented such that it projects radially inwardly on the loop.

20 10. The anastomotic device according to claim 1, wherein the at least one tissue-penetrating protrusion forms an acute angle with respect to a radial centerpoint of the loop.

25 11. The anastomotic device according to claim 1, further including at least one spacer element disposed on the flexible member between adjacent anchoring elements.

 12. The anastomotic device according to claim 11, wherein the at least one spacer element is slidably mounted to the flexible member.

30 13. The anastomosis device according to claim 11, wherein the at least one spacer element is a tubular member having an inner, axial opening that is configured and dimensioned to receive the flexible member.

35 14. The anastomotic device according to claim 11, wherein the at least one spacer element is flexible.

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5 15. The anastomotic device according to claim 1, wherein the anchoring element is a substantially tubular member having an axial opening formed therein that is configured and dimensioned to receive the flexible member.

10 16. An anastomotic device, comprising:

 a flexible member having a first end and a second end and first and second surfaces;

 a locking mechanism secured to one of the first or second ends, the locking mechanism having at least one opening therein for receiving the other of the first or second ends such that the locking mechanism facilitates the formation of a loop having an adjustable size; and

15 one or more anchoring elements mounted upon the flexible member, each anchoring element having at least one tissue penetrating protrusion formed on one surface thereof

20 17. An exovascular anastomotic device, comprising:

 a flexible member having first and second ends and first and second surfaces, the member having a locking mechanism formed on the first end thereof, the locking mechanism including an opening formed therein through which the second end of the flexible member is inserted to form a loop of an adjustable diameter;

25 a plurality of projections formed on one of the first or second surfaces of the flexible member, the projections cooperating with the locking mechanism to permit the second end of the flexible member to move relative to the locking mechanism in one direction to effect a decrease in the diameter of the loop;

30 one or more anchor members, each having a body defining an axial opening within which the flexible member is disposed to slidably mount the anchor members on the flexible member; and

 at least one tissue penetrating projection formed on an inwardly facing surface of each anchoring member.

5 18. The device according to claim 17, further comprising at least one spacer member slidably disposed on the flexible member between at least two adjacent anchoring members.

10 19. An anastomotic device for anastomosing an end of a first vessel to a side of a second vessel, comprising:

an anastomotic ring adapted to be disposed about an end of the first vessel;

a harness assembly including a harness ring having a radially overlapping portion with respect to the anastomotic ring;

15 at least one flexible member secured to the harness ring, the at least one flexible member forming a loop of adjustable size.

20 20. The anastomotic device according to claim 19, further including a locking mechanism mounted on the at least one flexible member.

21. The anastomotic device according to claim 19, wherein the anastomotic ring is tapered about a circumferential outer surface.

22. A method for conducting a sutureless anastomotic procedure, comprising the steps of:

25 providing a sutureless anastomotic device including a flexible member in the form of a loop having an adjustable diameter, at least one anchoring element having an inwardly projecting tissue penetrating member;

placing an end of a first vessel segment within the loop;

30 overlapping portions of the end of the first vessel segment and an end of a second vessel segment; and

applying an axial force to a first end of the flexible member to reduce the diameter of the loop such that the tissue penetrating members penetrate the overlapping portions of the first and second vessels to anastomose the first and second vessel.

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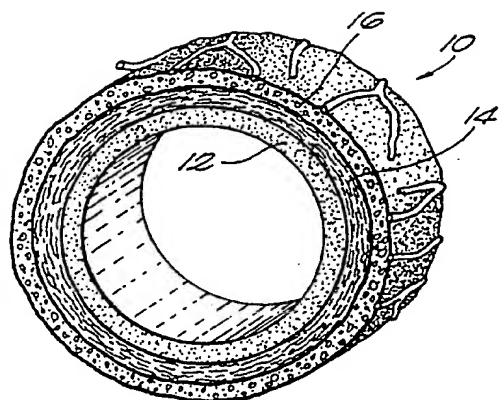


FIG. 1
(PRIOR ART)

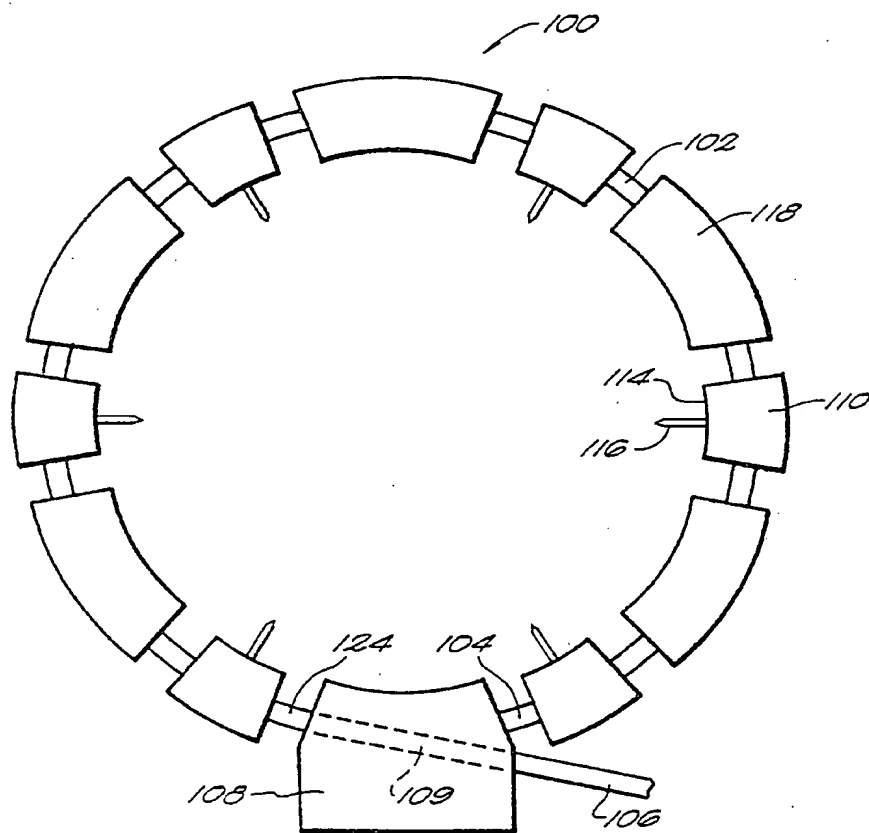


FIG. 2

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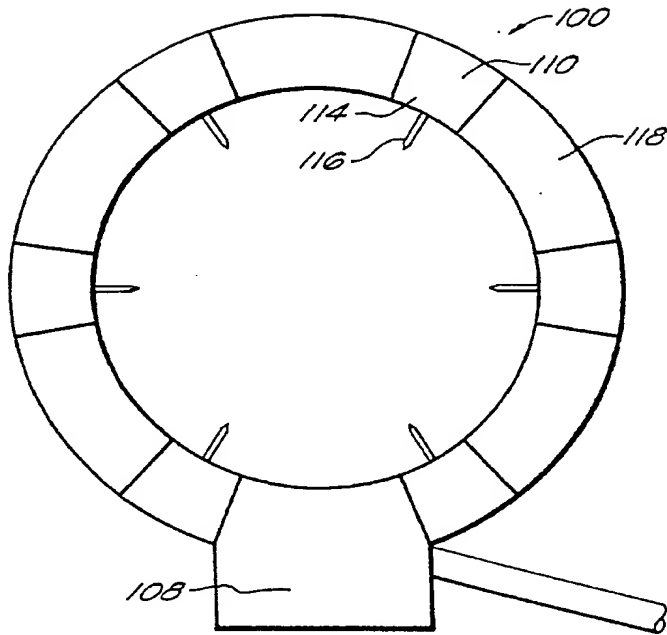


FIG. 3

FIG. 4

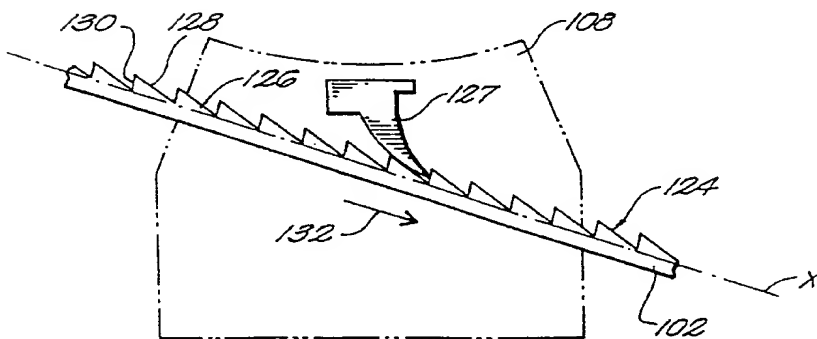
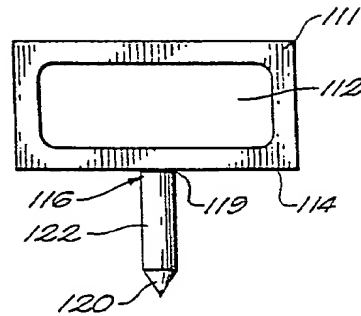
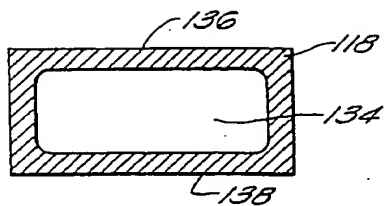
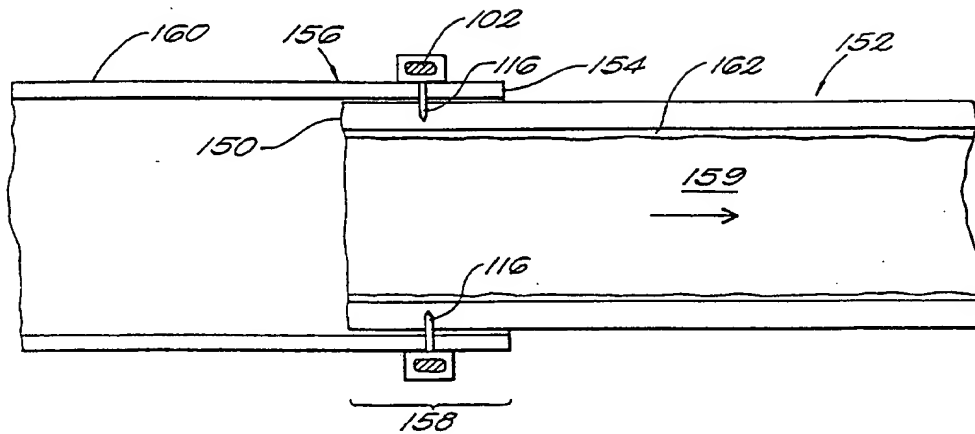
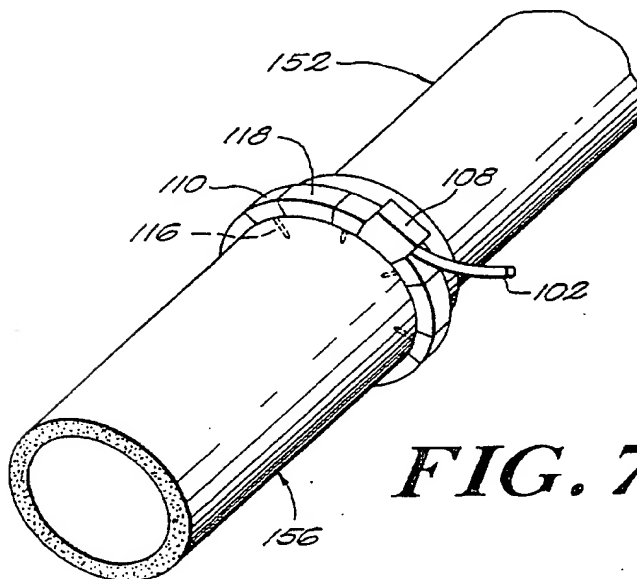
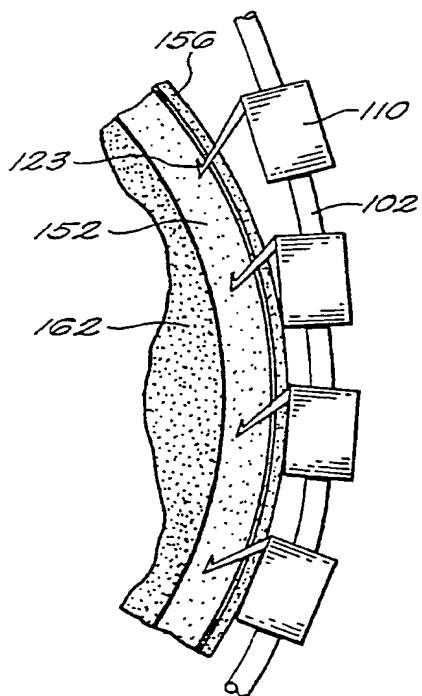
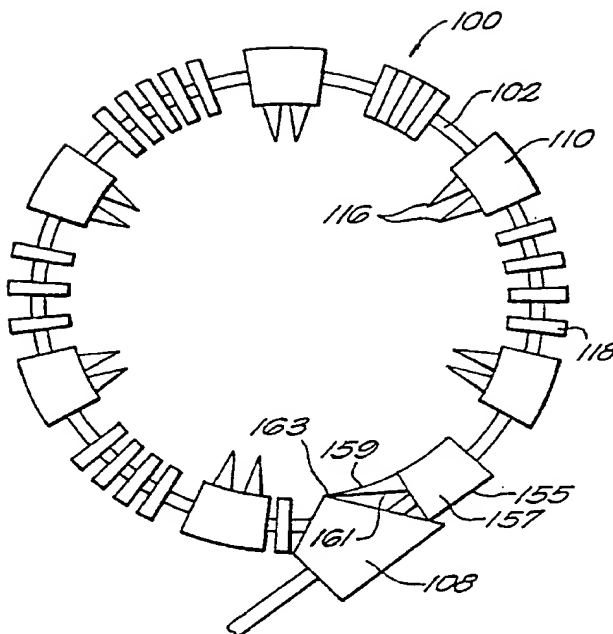
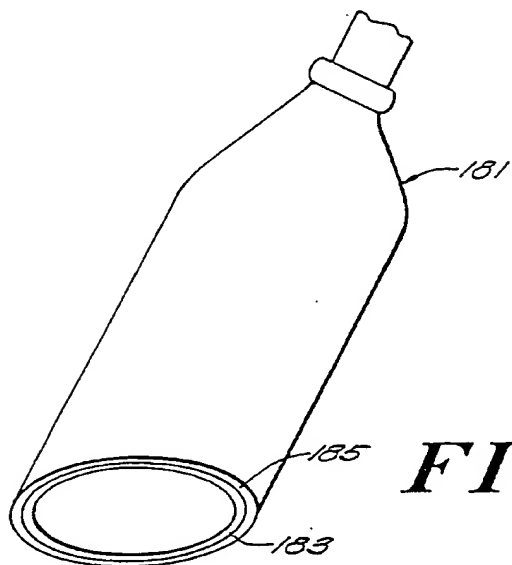


FIG. 5

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**FIG. 6****FIG. 7****FIG. 7A**

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**FIG. 7B****FIG. 7C****FIG. 7D**

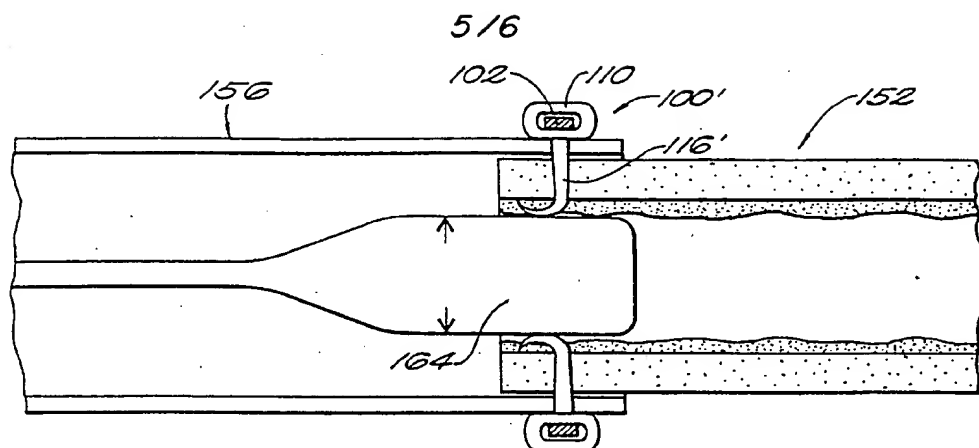


FIG. 8

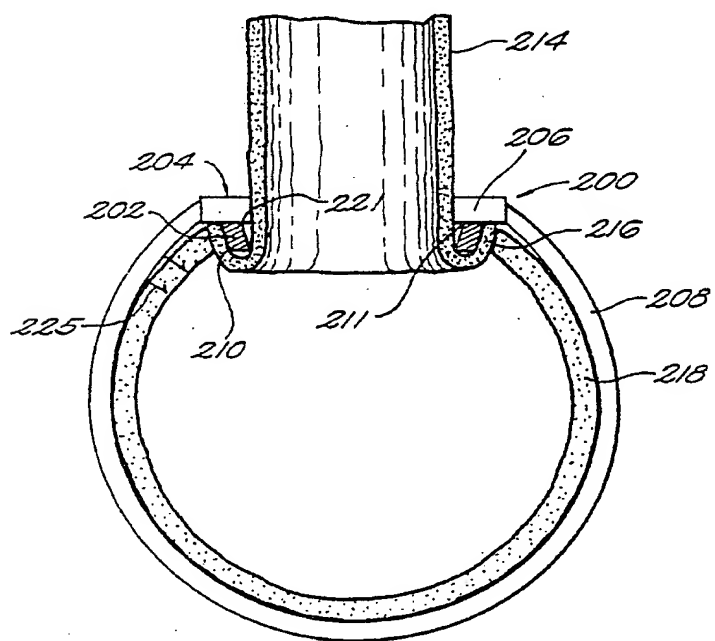


FIG. 9

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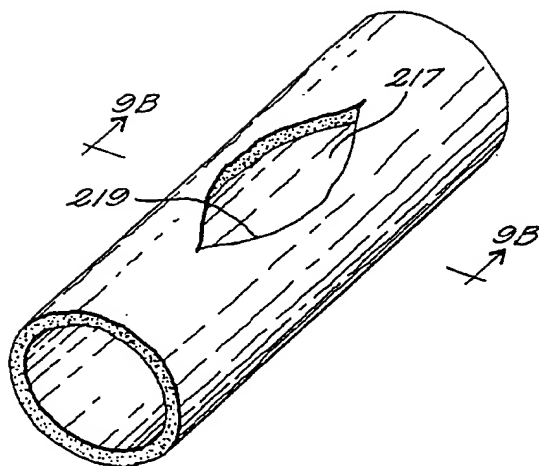


FIG. 9A

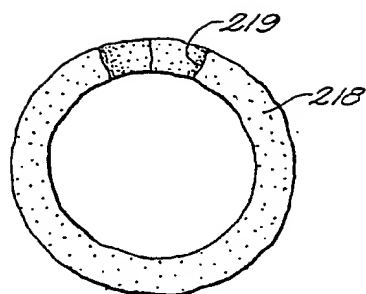


FIG. 9B

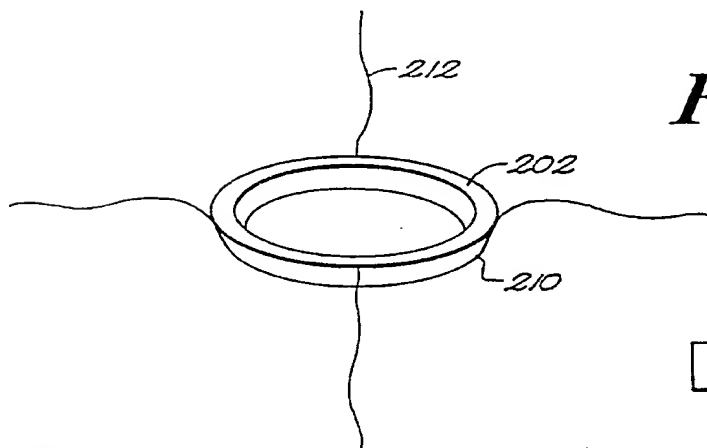


FIG. 10

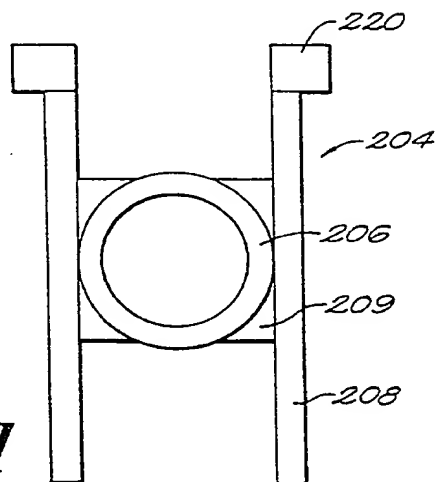


FIG. 11

INTERNATIONAL SEARCH REPORT

International application No.
PCT/US98/04387

A. CLASSIFICATION OF SUBJECT MATTER

IPC(6) : A61B 17/04
US CL : 606/151, 153

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

U.S. : 606/139, 151, 153

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practicable, search terms used)

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	US 4,693,249 A (SCHENCK et al) 15 September 1987, col. 2 lines 21-53.	1-22
A	US 5,234,447 A (KASTER et al) 10 August 1993, col. 7 lines 21-57.	1-22
A	US 5,486,187 A (SCHENCK) 23 January 1996, col. 3 lines 41-65.	1-22

☐ Further documents are listed in the continuation of Box C. ☐ See patent family annex.

* Special categories of cited documents:	*T* later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention
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L document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)	*Z* document member of the same patent family
O document referring to an oral disclosure, use, exhibition or other means	
P document published prior to the international filing date but later than the priority date claimed	

Date of the actual completion of the international search

05 JUNE 1998

Date of mailing of the international search report

29 JUN 1998

Name and mailing address of the ISA/US
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